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E3
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DIAZIDE-(2)-5-SULFONATE/CN
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E23
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=> D L1
     ANSWER 1 OF 1 REGISTRY COPYRIGHT 2007 ACS on STN
     700-13-0 REGISTRY
ED
     Entered STN: 16 Nov 1984
     1,4-Benzenediol, 2,3,5-trimethyl- (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Hydroquinone, trimethyl- (6CI, 7CI, 8CI)
OTHER NAMES:
     ψ-Cumohydroquinone
CN
     1,4-Dihydroxy-2,3,5-trimethylbenzene
CN
CN
     2,3,5-Trimethyl-1,4-benzenediol
CN
     2,3,5-Trimethyl-1,4-hydroquinone
CN
     2,3,5-Trimethyl-p-hydroquinone
    2,3,5-Trimethylhydroquinone
CN
CN
     NSC 401617
     Pseudocumohydroquinone
CN
CN
     Trimethyl-p-hydroquinone
     Trimethylhydroquinone
CN
MF
     C9 H12 O2
CI
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                  AGRICOLA, ANABSTR, BEILSTEIN*, BIOSIS, CA, CAOLD, CAPLUS,
LC
     STN Files:
       CASREACT, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, CSNB, DETHERM*,
       IFICDB, IFIPAT, IFIUDB, MEDLINE, MSDS-OHS, PROMT, PS, RTECS*, SPECINFO,
       SYNTHLINE, TOXCENTER, ULIDAT, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
                     EINECS**, NDSL**, TSCA**
     Other Sources:
         (**Enter CHEMLIST File for up-to-date regulatory information)
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Me OH Me
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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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923 REFERENCES IN FILE CA (1907 TO DATE)
6 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
925 REFERENCES IN FILE CAPLUS (1907 TO DATE)
31 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
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E3
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E18
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E19
                    2,3,5-TRIMETHYLPHENYL 1,2-NAPHTHOQUINONE
E20
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DIAZIDE-(2)-5-SULFONATE/CN
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E21
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E22
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E23
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E25
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E3
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E8
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E9
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E14
E15
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                    2,3,5-TRIMETHYLPHENANTHRENE/CN
E16
             1 ·
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E17
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E18
             1
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E19
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                   2,3,5-TRIMETHYLPHENYL 1,2-NAPHTHOQUINONE
E20
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                   2,3,5-TRIMETHYLPHENYL 2,4,5-TRIMETHYLPHENYL SULFONE/CN
E21
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                   2;3,5-TRIMETHYLPHENYL ACETATE/CN
E22
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E23 ·
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                   2,3,5-TRIMETHYLPHENYL ACETIC ACID/CN
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E25
                   2,3,5-TRIMETHYLPHENYL METHYLCARBAMATE/CN
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                                                  SINCE FILE
                                                                  TOTAL
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                                                                SESSION
FULL ESTIMATED COST
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=> S L1
           925 L1
L_2
=> S ALKONATE
             3 ALKONATE
             2 ALKONATES
            . 5 ALKONATE
                 (ALKONATE OR ALKONATES)
=> S DIALKONATE
L4
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=> S KETOISOPHORONE
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             1 KETOISOPHORONES
L5
            82 KETOISOPHORONE
                 (KETOISOPHORONE OR KETOISOPHORONES)
=> S L2 AND L5
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L6 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN AN 2005:570864 CAPLUS

3 L2 AND L5

DN 143:77878

=> D BIB ABS HITSTR

```
TI
     Catalytic esterification process for the manufacture of
     trimethylhydroquine dialkanoates
IN
     Bonrath, Werner; Foricher, Yann
PA
     DSM IP Assets B. V., Neth.
     PCT Int. Appl., 12 pp.
SO
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 1
     PATENT NO.
                                DATE
                                             APPLICATION NO.
                                                                    DATE
                         KIND
                                             -----
                         _ _ _ _
PΤ
     WO 2005058792
                         · A1
                                20050630
                                             WO 2004-EP13903
                                                                    20041207
             AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
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             LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI,
             NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
             TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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                                20060830
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                                20.061227
                                             CN 2004-80035561
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     JP 2007516262
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                                20070621
                                             JP 2006-544280
                                                                    20041207
     US 2007049762
                                 20070301
                                             US 2006-582672
                                                                    20060629
                          A1
PRAI EP 2003-28811
                                20031215
                          Α
     WO 2004-EP13903
                          W
                                20041207
OS
     CASREACT 143:77878
     A process for the manufacture of a 2,3,5-trimethylhydroquinone dialkanoate
AB.
     comprises reacting ketoisophorone with an acylating agent in the
     presence of an indium salt as the catalyst. Preferred are indium(III)
     salts such as indium trichloride or indium tris(trifluoromethanesulfonate)
       Process for the manufacture of 2,3,5-trimethylhydroquinone using
     2,3,5-trimethylhydroquinone dialkanoate as the starting material, especially a
     process for the manufacture of 2,3,5-trimethylhydroquinone by
     transesterification of 2,3,5-trimethylhydroquinone dialkanoate, as well as
     a process for the manufacture of \alpha-tocopherol and its alkanoates, especially
of
     (all-racemic)-\alpha-tocopherol and its acetate, comprising the reaction
     of ketoisophorone to 2,3,5-trimethylhydroquinone dialkanoate are
     presented.
     700-13-0, 2,3,5-Trimethylhydroquinone
IT
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (esterification of)
RN
     700-13-0 CAPLUS
CN
     1,4-Benzenediol, 2,3,5-trimethyl- (CA INDEX NAME)
Me.
           OH
```

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> D BIB ABS HITSTR 2-3

L6 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2001:740313 CAPLUS

DN 136:71494

TI Industrial application of Nafion-systems in rearrangement-aromatisation, transesterification, alkylation, and ring-closure reactions

AU Schneider, M.; Zimmermann, K.; Aquino, F.; Bonrath, W.

CS Vitamins and Fine Chemicals Division, Chemical Process Technology, F. Hoffmann-La Roche Ltd., Basel, CH-4070, Switz.

SO Applied Catalysis, A: General (2001), 220(1-2), 51-58 CODEN: ACAGE4; ISSN: 0926-860X

PB Elsevier Science B.V.

DT Journal

LA English

Nafion, a perfluorinated sulfonic acid ion-exchange polymer, is known to AB be a very strong Bronsted acid. Thus, Nafion NR 50 and Nafion/SiO2 with 15 weight % Nafion-loading were selected, in order to elucidate the potential for rearrangement-aromatization of ketoisophorone (KIP) to 2,3,5-trimethylhydroquinone diacetate (TMHQ-DA) in presence of acetic acid anhydride as acylating agent, transesterification of TMHQ-DA to 2,3,6-trimethylhydroguinone monoacetate (TMHQ-1-MA) and reaction of isophytol (IP) with trimethylhydroquinone (TMHQ) to (all-rac)- α tocopherol. For the rearrangement-aromatization of KIP to TMHQ-DA supported Nafion/SiO2 was markedly more active than the unsupported Nafion NR 50. Both Nafion-systems generally revealed remarkably high selectivity, which ranged up to 94 GC-a% TMHQ-DA at high conversion. major side-product was 3,4,5-trimethylcatechol diacetate. In case of Nafion/SiO2, pre-treatment under vacuum and especially grinding of the extrudates seemed to increase activity at comparably high selectivity. Recycling Nafion/SiO2 after filtering and rinsing with acetic acid anhydride led to gradually decreasing activity. Without intermediate isolation of TMHQ-DA, Nafion/SiO2 was active and selective for the formation of TMHQ-1-MA. Using the Nafion systems in the reaction of IP and TMHQ revealed that remarkably high conversion of IP (>95%) and. compared with zinc chloride/Bronsted acid or BF3-catalyzed reaction good yields (≈92%) and selectivities were obtained. We found a strong dependency on the solvent polarity. In further expts., the recovery of the catalyst was tested.

IT 700-13-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(Nafion-catalyzed preparation and cyclization reaction of trimethylhydroquinone with isophytol)

RN 700-13-0 CAPLUS

CN 1,4-Benzenediol, 2,3,5-trimethyl- (CA INDEX NAME)

RE.CNT 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2007 ACS on STN

AN 2001:297544 CAPLUS

DN 134:295967

TI Process for the preparation of esters of chroman derivatives

IN Weigel, Horst; Krill, Steffen; Hasselbach, Hans Joachim; Huthmacher, Klaus

PA Degussa-Huls A.-G., Germany

SO Eur. Pat. Appl., 14 pp.

CODEN: EPXXDW

DT Patent

LA German

FAN.CNT 1

		PATENT NO.					KINI	D DATE	DATE		APPLICATION NO.				DATE		
																'	
3	ΡI	EP 1094062				A1 20010425			EP 2000-116383				20000728				
			R:	AT,	BE,	CH,	DE,	DK, ES,	FR,	GB, GF	?, IT,	LI,	LU,	NL,	SE,	MC,	PT,
				ΙE,	SI,	LT,	LV,	FI, RO									
		DE 19951006				A1	2001	0426	DE	1999-	19951	.006		19	9991	22	
		IN	1887	10	•		A1	2002	1026	IN	2000-0	CA582	?		20	0001	16
		CA	2323	840			A1	2001	0422	· CA	2000-2	23238	340		20	0001	19
		CN	1308	077			A	2001	0815	CN	2000-	12987	0		. 20	0001	19
		US	6329	535			B1	2001	1211	US	2000-	69231	.3		20	0001	20
		IL	1391	78			. A	2004	0725	IL	2000-3	13917	78		20	0001	020
]	PRAI	DE	1999	-1999	51006	5	A	1999	1022								
		·															

OS CASREACT 134:295967

AB Process for the preparation of chroman ester derivs., in that one, (1.1) reacts tech. pure ketoisophorone with a acylating agent in the presence of a protonic acid to give a trimethylhydroquinone ester, and (1.2) this ester is reacted with allyl alc. or its derivative in the presence of zinc halide and a proton cleaving acid, and (1.3) carried out in the presence of an organic solvent, is characterized by (1.1.1) the solution in reaction

step

(1.1) is cooled to a temperature between 5 and 40°, (1.1.2) the crystallized product is separated and washed, (1.1.3) that the saved filtrate from the separated product and the washes is out into the solvent for future reaction step (1.1), (1.2.1) the washed product without drying is introduced into reaction (1.2) and, (1.2.2) the desired product isolated after further acylation. Thus, ketoisophorone is reacted with Ac2O in the presence of triflic acid, to give 87% trimethylhydroquinone diacetate; the latter is reacted with isophytol in PhMe containing ZnBr2 and HBr followed by Ac2O to give 95.6% vitamin E acetate.

IT 700-13-0DP, Trimethylhydroquinone, esters

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of chroman ester derivs. via acylation of ketoiosphorone followed by reaction with allyl alc. derivs.)

RN 700-13-0 CAPLUS

CN 1,4-Benzenediol, 2,3,5-trimethyl- (CA INDEX NAME)

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT